

**K022579 STRYKER NAVIGATION SYSTEM-KNEE MODULE**Jun 2, 2003  
301 days to decisionK022579 · Product code: **HAW** · Neurology  
Source: <https://www.510kdatabase.net/k022579/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Neurological Stereotaxic Instrument (HAW)
Date received	Aug 5, 2002
Decision date	Jun 2, 2003
Days to decision	301 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Stryker Leibinger</b>
Location	Kalamazoo, MI, US
Contact	KELLI J BITTERBURG
510(k) history	12 submissions · 12 cleared · 1999-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k022579/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 19, 2026